## FOOD AND DRUG ADMINISTRATION (FDA)

Center for Drug Evaluation and Research (CDER)

Meeting of the Pulmonary-Allergy Drugs Advisory Committee (PADAC)

FDA White Oak Campus, Building 31, the Great Room, White Oak Conference Center (Rm. 1503), Silver Spring, MD

January 30, 2013

## **DRAFT AGENDA**

The committee will discuss the New Drug Application (NDA) for mannitol inhalation powder (Bronchitol), sponsored by Pharmaxis, for the management of cystic fibrosis (CF) in patients aged 6 years and older to improve pulmonary function.

Call to Order 8:00 a.m. Introduction of Committee Chairperson, Pulmonary-Allergy Drugs Advisory Committee (PADAC) 8:05 a.m. Conflict of Interest Statement Cindy Hong, PharmD Designated Federal Officer, PADAC 8:10 a.m. **Opening Remarks** Anthony Durmowicz, MD Clinical Team Leader, Division of Pulmonary, Allergy, and Rheumatology Products (DPARP), Office of Drug Evaluation II (ODE-II), Office of New Drugs (OND), CDER, FDA 8:15 a.m. **Sponsor Presentations Pharmaxis** Ronald Dundore, PhD Introduction VP, US Regulatory Affairs Pharmaxis Unmet Medical Need Felix Ratjen, MD, PhD Professor, Respiratory Medicine University of Toronto Hospital for Sick Children Efficacy Howard Fox, MD Chief Medical Officer **Pharmaxis** Safety Brett Charlton, MD, PhD **Medical Director Pharmaxis** Risk/Benefit and Clinical Perspective Patrick Flume, MD Professor, Pulmonary and Critical Care Medicine Medical University of South Carolina

Clarifying Questions to the Presenters

9:30 a.m.

## FOOD AND DRUG ADMINISTRATION (FDA)

Center for Drug Evaluation and Research (CDER)

Meeting of the Pulmonary-Allergy Drugs Advisory Committee (PADAC)

FDA White Oak Campus, Building 31, the Great Room, White Oak Conference Center (Rm. 1503), Silver Spring, MD

January 30, 2013

## **DRAFT AGENDA (cont.)**

9:45 a.m. **BREAK** 

10:00 a.m. **FDA Presentations** 

Overview of the Clinical Program Kimberly Witzmann, MD

Clinical Reviewer

DPARP, ODE-II, CDER, FDA

Statistical Review of Efficacy Feng Zhou, MS

Statistical Reviewer

Division of Biostatistics II (DB-II)

Office of Biostatistics (OB)

Office of Translational Sciences (OTS), CDER, FDA

**Thomas Permutt, PhD** 

Director, Division of Biostatistics II (DB-II)

Office of Biostatistics (OB)

Office of Translational Sciences (OTS), CDER, FDA

Clinical Review of Efficacy, Safety,

and Risk/Benefit

Kimberly Witzmann, MD

11:15 a.m. Clarifying Questions to the Presenters

11:30 a.m. **LUNCH** 

12:30 p.m. Open Public Hearing

1:30 p.m. Charge to the Committee Anthony Durmowicz, MD

1:40 p.m. Questions to the Committee and Committee Discussion

2:30 p.m. **BREAK** 

2:45 p.m. Questions to the Committee and Committee Discussion (cont.)

4:00 p.m. **ADJOURNMENT** 

Page 2 of 2